Food and Drug Administration, HHS

§522.1890 Sterile prednisone suspension.

- (a) [Reserved]
- (b)(1) *Specifications*. Each milliliter of sterile aqueous suspension contains 10 to 40 milligrams of prednisone.
- (2) *Sponsor*. See 000061 in §510.600(c) of this chapter.
- (3) Conditions of use—(i) Amount. Administer intramuscularly as follows:
- (a) Horses. 100 to 400 milligrams, repeating if necessary. If no response is observed after 3 to 4 days of therapy, reevaluate diagnosis.¹
- (b) Dogs and cats. 0.25 to 1.0 milligram per pound of body weight for 3 to 5 days or until a response is noted. Treatment may be continued with an orally administered dose.¹
- (ii) *Indications for use*. It is used for conditions requiring an anti-inflammatory agent.¹
- (iii) Limitations. 1 Do not use in viral infections. Except in emergency therapy, do not use in animals with tubernephritis, culosis. chronic orCushings's disease. With infections, use appropriate antibacterial therapy with and for at least 3 days after discontinuance of use and disappearance of all signs of infection. Clinical and experimental data have demonstrated that corticosteroids administered orally or parenterally to animals may induce the first stage of parturition when administered during the last trimester of pregnancy and may precipitate premature parturition followed hv dystocia, fetal death, retained placenta, and metritis. Not for use in horses intended for food. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

[45 FR 13446, Feb. 29, 1980, as amended at 50 FR 6160, Feb. 14, 1985; 52 FR 7832, Mar. 13, 1987]

§522.1920 Prochlorperazine, isopropamide for injection.

(a) Specifications. Prochlorperazine, isopropamide for injection, veterinary, contains in each milliliter, 6 milli-

grams of prochlorperazine edisylate (equivalent to 4 milligrams prochlorperazine), and 0.38 milligrams of isopropamide iodide (equivalent to 0.28 milligrams of isopropamide) in buffered aqueous solution.

- (b) *Sponsor*. See No. 000069 in §510.600(c) of this chapter.
- (c) Conditions of use. (1) The drug is used in dogs and cats in which gastro-intestinal disturbances are associated with emotional stress.
- (2) Dosage is administered by subcutaneous injection twice daily as follows:

Weight of animal in pounds	Dosage in Milliliters
Up to 4	0.25
5 to 14	0.5-1
15 to 30	2-3
30 to 45	3-4
45 to 60	4–5
Over 60	6

Following the last injection, administer prochlorperazine and isopropamide sustained release capsules as indicated.

(3) For use only by or on the order of a licensed veterinarian.

[40 FR 13858, Mar. 27, 1975, as amended at 56 FR 50653, Oct. 8, 1991; 60 FR 55659, Nov. 2, 1995]

§ 522.1940 Progesterone and estradiol benzoate in combination.

- (a) [Reserved]
- (b) Sponsors. See 000856 in $\S510.600(c)$ of this chapter for use as in paragraphs (d)(1)(i)(A), (d)(1)(ii), (d)(2)(iii), (d)(2)(iii), and (d)(3) of this section. See 021641 in $\S510.600(c)$ of this chapter for use as in paragraphs (d)(1) and (d)(2)(i) through (d)(2)(iii)(A) of this section.
- (c) Related tolerances. See §§ 556.240 and 556.540 of this chapter.
- (d) Conditions of use. It is used for implantation in animals as follows:
- (1) Suckling beef calves—(i) Amount. (A) 100 milligrams of progesterone and 10 milligrams of estradiol benzoate in four pellets per implant dose.
- (B) 100 milligrams of progesterone and 10 milligrams of estradiol benzoate in four pellets with 29 milligrams of tylosin tartrate as a local antibacterial in one pellet per implant dose.

¹These conditions are NAS/NRC reviewed and deemed effective. Applications for these uses need not include effectiveness data as specified by §514.111 of this chapter, but may require bioequivalency and safety information.